

K132066 INFLAMMA DRYNov 21, 2013
141 days to decisionK132066 · Product code: **PFQ** · Chemistry
Source: <https://www.510kdatabase.net/k132066/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Mmp-9 Test System (PFQ)
Date received	Jul 3, 2013
Decision date	Nov 21, 2013
Days to decision	141 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Rapid Pathogen Screening, Inc.
Location	Sarasota, FL, US
Contact	Douglas Bueschel
510(k) history	2 submissions · 2 cleared · 2011-2013

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k132066/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated June 8, 2026